# ALL DAY ALLERGY RELIEF- cetirizine hcl capsule QUALITY CHOICE (Chain Drug Marketing Association)

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# **Cetirizine Hydrochloride Tablets**

# Active Ingredient (in each capsule)

Cetirizine HCl 10 mg

# **Purpose**

**Antihistamine** 

#### Uses

temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

- runny nose
- sneezing
- itchy, watery eyes
- itching of the nose or throat

# **Warning**

#### Do not use

if you have ever had an allergic reaction to this product or any of its ingredients or to an antihistamine containing hydroxyzine.

# Ask a doctor before use if you have

liver or kidney disease. Your doctor should determine if you need a different dose.

# Ask a doctor or pharmacist before use if you are

taking tranquilizers or sedatives.

# When using this product

- drowsiness may occur
- alcohol, sedatives, and tranquilizers may increase drowsiness
- avoid alcoholic drinks
- be careful when driving a motor vehicle or operating machinery

## Stop use and ask a doctor if

an allergic reaction to this product occurs. Seek medical help right away.

# If pregnant or breast-feeding

- if breast-feeding: not recommended
- if pregnant: ask a health professional before use.

## Keep out of reach of children

In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

#### **Directions**

Adults and children 6 years and over	One 10 mg capsule once daily; do not take more than one 10 mg capsule in 24 hours. A 5 mg product may be appropriate for less severe symptoms.
Adults 65 years and over	ask a doctor.
Children under 6 years of age	ask a doctor
Consumers with liver or kidney disease	ask a doctor

#### Other Information

- store at 20° to 25°C (68° to 77° F)
- avoid high humidity and excessive heat above 40°C (104°F)
- protect from light

# **Inactive Ingredients**

Butylated hydroxytolene, gelatin, glycerin, mannitol, pharmaceutical ink, polyethylene glycol 400, purified water, sodium hydroxide, Sorbitan, Sorbitol

## Questions or comments?

Call 1-248-449-9300 Monday-Friday 9AM-5PM EST

# **Principal Display Panel**

\*Compare to the Active Ingredient in ZYRTEC®

All Day Allergy Relief

Indoor & Outdoor Allergies

Cetirizine HCI Capsules, 10 mg

**Antihistamine** 

24 HOUR RELIEF OF:

Sneezing

Runny Nose

Itchy, Watery Eyes

Itchy Throat or Nose

Softgels\*\*

(\*\*Liquid-Filled Capsules)

\*This product is not manufactured or distributed by McNeil Consumer Healthcare, distributor of Zyrtec®.

# TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.

## KEEP OUTER CARD FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

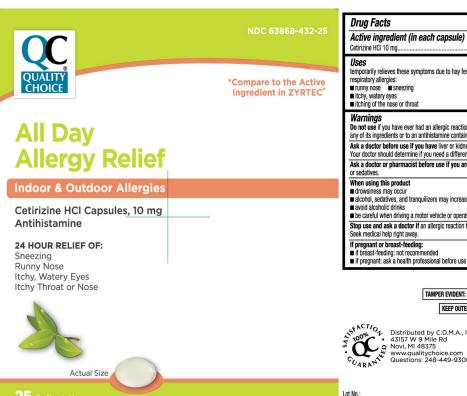
Distributed by: C.D.M.A., Inc.©

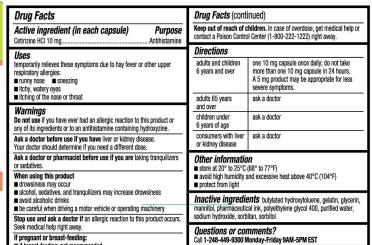
43157 W 9 Mile Rd

Novi, MI 48375

www.qualitychoice.com

## **Product Label**





TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.

KEEP OUTER CARD FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

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\*This product is not manufactured or distributed by McNeil Consumer

Healthcare, distributor of Zyrtec®

Exp. Date:

## **QUALITY CHOICE All Day Allergy Relief**

# **ALL DAY ALLERGY RELIEF**

cetirizine hcl capsule

#### **Product Information**

HUMAN OTC DRUG NDC:63868-432 **Product Type Item Code (Source)** 

**Route of Administration ORAL** 

# **Active Ingredient/Active Moiety**

**Ingredient Name Basis of Strength** Strength CETIRIZINE HYDROCHLORIDE (UNII: 640047KTOA) (CETIRIZINE -**CETIRIZINE** 10 mg UNII:YO7261ME24) **HYDROCHLORIDE** 

Inactive Ingredients	
Ingredient Name	Strength
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
MANNITOL (UNII: 3OWL53L36A)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SORBITAN (UNII: 6092ICV9RU)	
SORBITOL (UNII: 506T60A25R)	

## BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)

Product Characteristics					
Color	yellow	Score	no score		
Shape	OVAL	Size	13mm		
Flavor		Imprint Code	10		
Contains					

P	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:63868-432- 25	1 in 1 BOX	06/30/2018	02/23/2024			
1		25 in 1 BOTTLE; Type 0: Not a Combination Product					

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
ANDA	ANDA207235	06/30/2018	02/23/2024	

Labeler - QUALITY CHOICE (Chain Drug Marketing Association) (011920774)

Revised: 11/2021 QUALITY CHOICE (Chain Drug Marketing Association)